

K990665 BQ/BM INTEGRATED LASER DELIVERY SYSTEMMay 17, 1999
76 days to decisionK990665 · Product code: **HQF** · Ophthalmic
Source: <https://www.510kdatabase.net/k990665/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Laser, Ophthalmic (HQF)
Date received	Mar 2, 1999
Decision date	May 17, 1999
Days to decision	76 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	American Laser Corp.
Location	Mchenry, IL, US
Contact	DANIEL HOEFER
510(k) history	5 submissions · 5 cleared · 1982-2001

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k990665/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 21, 2026